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Supplementary Information

Chondroinductive Factor-Free Chondrogenic Differentiation of Human Mesenchymal Stem Cells in

Graphene Oxide-Incorporated Hydrogels

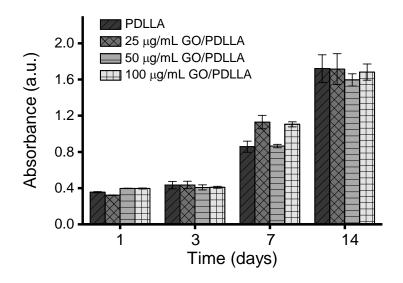


Figure S1. Cellular metabolism of hBMSCs encapsulated in PDLLA hydrogel with or without the presence of GO. After culturing in growth medium for 1, 3, 7 and 14 days, results from MTS assay showed no differences in cellular metabolism between control (no GO) and GO-added constructs. Valued are mean \pm S.D., n=3.

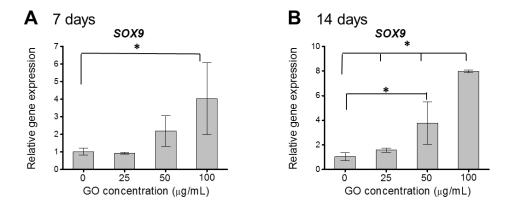


Figure S2. SOX 9 expression in hBMSCs seeded in hydrogel cultures as a function of GO incorporation. hBMSC (20×10^6 cells/mL)-laden PDLLA hydrogels containing 0, 25, 50 and 100 µg/mL GO were cultured for (A) 7 days and (B) 14 days, and analyzed for SOX9 by real-time PCR. Values are mean \pm S.D. (n=3), expressed relative to the level in the absence of GO. The results showed significant SOX9 up-regulation upon exposure to GO, expecially at 100 µg/mL. *, p<0.05

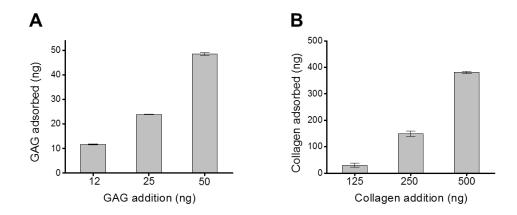


Figure S3. (A) Sulfated GAG and (B) collagen adsorption on GO nanosheets. The loading capacity of the GO nanosheet is indicated by the increased GAG and collagen adsorbed as a function of increasing amounts added. Values are mean \pm S.D.